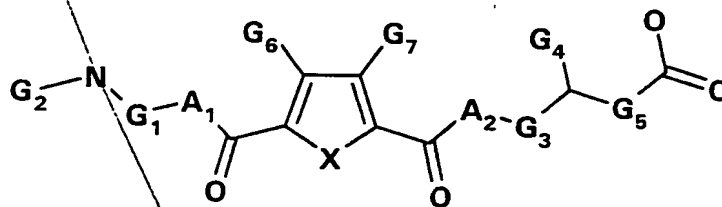


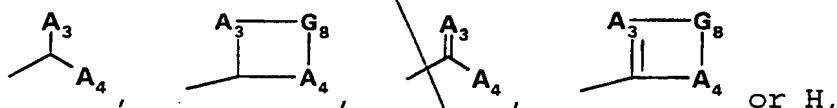
We claim:

1. A compound of the formula:



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wherein X is selected from the group comprising O and S;
 wherein A₁ and A₂ are individually selected from the
 group comprising O, S and N;
 wherein G₁ and G₃ are C₁₋₄ alkyl chains;
 wherein G₅ is a C₀₋₄ alkyl chain;
 wherein G₂ is selected from the group comprising:



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wherein A₃ and A₄ are individually selected from the
 group comprising O, N, or S, and G₈ is a C₁₋₄ alkyl
 chain;

wherein G₄ is a C₅₋₈ aryl, a C₅₋₈ arylsulfonylamino, an
 C₅₋₈ arylamino; and

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wherein G₆ and G₇ are individually selected from the
 group comprising H, F, Cl, I, Br and a C₁₋₄ alkyl.

2. The compound of claim 1, wherein X is S.

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3. The compound of claim 1, wherein X is O.

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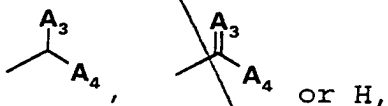
4. The compound of claim 1, wherein A₁ is N.

5. The compound of claim 1, wherein A₁ is O.

Sub
B3

6. The compound of claim 1, wherein A_2 is N.
7. The compound of claim 1, wherein A_2 is O.
8. The compound of claim 1, wherein G_1 is a C_1 alkyl.
9. The compound of claim 1, wherein G_1 is $-(CH_2)_0-$.
10. The compound of claim 1, wherein G_1 is a C_2 alkyl.
11. The compound of claim 1, wherein G_1 is a C_3 alkyl.
12. The compound of claim 1, wherein G_3 is a C_1 alkyl.
13. The compound of claim 1, wherein G_3 is a C_2 alkyl.
14. The compound of claim 1, wherein G_5 is a C_1 alkyl.
15. The compound of claim 1, wherein G_5 is a C_2 alkyl.
16. The compound of claim 1, wherein G_2 is represented by the formula:

Sub
25 B4



wherein A_3 is selected from the group comprising O, S and N and A_4 is N.

17. The compound of claim 1, wherein G_2 is represented by the formula:



wherein A_3 and A_4 are individually selected from the group comprising N or O and G_8 is a C_{2-3} alkyl chain.

18. The compound of claim 1, wherein -N-G₂ forms a guanidino containing moiety.

5 19. The compound of claim 1, wherein -N-G₂ forms a urea containing moiety.

20. The compound of ~~claim~~ 1, wherein -N-G₂ forms a cyclic guanidino containing moiety.

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21. The compound of ~~claim~~ 1, wherein -N-G₂ forms a cyclic urea containing moiety.

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22. The compound of claim 1, wherein G₄ is phenylsulfonylamino.

23. The compound of claim 1, wherein G₄ is phenyl.

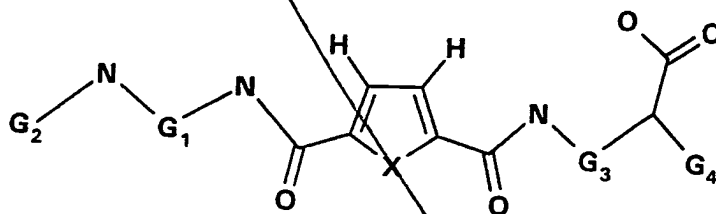
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24. The compound of claim 1, wherein G₆ and G₇ are halogens.

25. The compound of claim 1, wherein G₆ and G₇ are the same.

25 26. The compound of claim 1, wherein G₆ or G₇ are F.

27. The compound of claim 1 further represented by the formula:



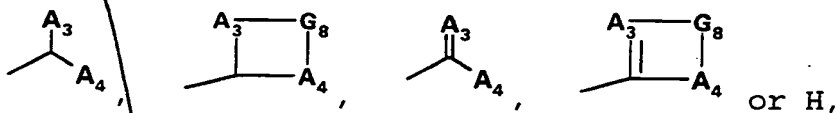
30 wherein X is selected from the group comprising O and S;

A_1 and A_2 are individually selected from the group comprising O, S and N;

G₁ and G₃ are C₁₋₄ alkyl chains;

G₂ is selected from the group comprising:

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wherein A₃ and A₄ are individually selected from the group comprising O, N, or S, and G₈ is a C₁₋₄ alkyl chain;

wherein G₄ is a C₅₋₈ aryl, a C₅₋₈ arylsulfonylamino, or a C₅₋₈ arylamino; and

wherein G_6 and G_7 are individually selected from the group comprising H, F, Cl, I, Br and a C_{1-4} alkyl.

28. The compound of claim 26, wherein X is S.

29. The compound of claim 26, wherein X is O.

20 30. The compound of claim 26, wherein G_1 is a C_1 alkyl.

31. The compound of claim 26, wherein G₁ is a C₂ alkyl.

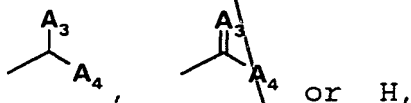
32. The compound of claim 26, wherein G₃ is a C₁ alkyl.

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33. The compound of claim 26, wherein G₃ is a C₂ alkyl.

34. The compound of claim 26, wherein G₂ is represented by the formula:

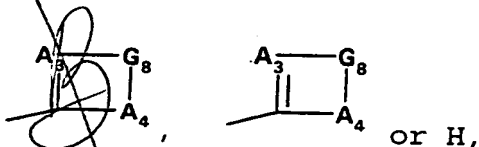
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wherein A_3 is selected from the group comprising 0, S and N and A_4 is N.

35. The compound of claim 26, wherein G_2 is represented by the formula:

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wherein A_3 and A_4 are individually selected from the group comprising N or O and G_8 is a C_{2-3} alkyl chain.

10 36. The compound of claim 26, wherein $-N-G_2$ forms a guanidino containing moiety.

15 37. The compound of claim 26, wherein $-N-G_2$ forms a urea containing moiety.

38. The compound of claim 26, wherein $-N-G_2$ forms is a cyclic guanidino containing moiety.

20 39. The compound of claim 26, wherein $-N-G_2$ forms a cyclic urea containing moiety.

40. The compound of claim 26, wherein G_4 is phenylsulfonylamino.

25 41. The compound of claim 26, wherein G_4 is phenyl.

42. A method of treating cancer comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.

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43. A method of treating a tumor comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.

44. A method of treating a solid tumor comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 5 45. A method of treating metastasis comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 10 46. A method of inhibiting angiogenesis comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 15 47. A method of inhibiting fibronectin binding comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 20 48. A method of inhibiting osteopontin binding comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 25 49. A method of treating foot and mouth disease comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 30 50. A method of treating osteoporosis comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 35 51. A method of treating restenosis comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
52. A method of treating ocular diseases comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.

53. A method of treating heart diseases comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 5 54. A method of treating arthritis comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 10 55. A method of treating diseases in which abnormal neovascularization occurs comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 15 56. A method of inhibiting α_v integrins comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 20 57. A method of inhibiting $\alpha_v\beta_3$ integrin comprising administering a pharmaceutically effective amount of the compound of claim 1 to a patient.
- 25 58. A pharmaceutical composition for treating cancer comprising a pharmaceutically effective amount of a compound of claim 1.
59. A pharmaceutical composition for treating tumor comprising a pharmaceutically effective amount of a compound of claim 1.
- 30 60. A pharmaceutical composition for treating solid tumor comprising a pharmaceutically effective amount of a compound of claim 1.
- 35 61. A pharmaceutical composition for treating metastasis comprising a pharmaceutically effective amount of a compound of claim 1.

62. A pharmaceutical composition for inhibiting angiogenesis comprising a pharmaceutically effective amount of a compound of claim 1.
- 5 63. A pharmaceutical composition for inhibiting fibronectin binding comprising a pharmaceutically effective amount of a compound of claim 1.
- 10 64. A pharmaceutical composition for inhibiting osteopontin binding comprising a pharmaceutically effective amount of a compound of claim 1.
- 15 65. A pharmaceutical composition for treating foot and mouth disease comprising a pharmaceutically effective amount of a compound of claim 1.
- 20 66. A pharmaceutical composition for treating osteoporosis comprising a pharmaceutically effective amount of a compound of claim 1.
- 25 67. A pharmaceutical composition for treating restenosis comprising a pharmaceutically effective amount of a compound of claim 1.
- 30 68. A pharmaceutical composition for treating ocular diseases comprising a pharmaceutically effective amount of a compound of claim 1.
- 35 69. A pharmaceutical composition for treating heart diseases comprising a pharmaceutically effective amount of a compound of claim 1.
70. A pharmaceutical composition for treating arthritis comprising a pharmaceutically effective amount of a compound of claim 1.

71. A pharmaceutical composition for treating diseases in which abnormal neovascularization occurs comprising a pharmaceutically effective amount of a compound of claim 1.

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72. A pharmaceutical composition for inhibiting α_v integrins comprising a pharmaceutically effective amount of a compound of claim 1.

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73. A pharmaceutical composition for inhibiting $\alpha_v\beta_3$ integrin comprising a pharmaceutically effective amount of a compound of claim 1.

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74. A combination useful for the treatment of cancer comprising at least one compound of claim 1 with at least one other anticancer agent or antiangiogenic agent.

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75. A combination useful for the treatment of cancer comprising at least one compound of claim 1 with at least one other anticancer agent selected from the group consisting of alkylating agents, antitumor antibiotics, antimetabolites, biological agents, hormonal agents, nitrogen mustard derivatives and plant alkaloids.

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